AMENDMENTS TO THE SPECIFICATION

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Please amend the paragraph beginning at line 18, through page 39, line 12 as indicated below:

In addition, regarding liver, as a sample for primary culture, a cell was isolated as follows. A sample was collected under a sterile procedure, the sample was suitably cut into strips, and recovered in a 50-ml conical tube. Twenty milliliters of 1% trypsin/EDTA/phosphate buffered saline was added thereto and suspended well, and culturing was performed at 37°C for 10 minutes while shaking. Two-hundred-microliters of a DNase I solution [solution in which 10 mg of DNase I was dissolved in 1 ml 0.15 M NaCl] was added to the resulting product, and 100 ml of 1 M MgCl2 was further added thereto, followed by culturing at 37°C for 10 minutes. Thereafter, the product was filtered with a nylon mesh having a diameter of 70 mm µm (manufactured by FALCON, trade name: cell strainer), and a D10 medium was added in the same amount as that of the product. Thereafter, the mixture was centrifuged at 1500 rpm for 5 minutes, and the supernatant was aspirated. Here, when a pellet was red due to contamination of erythrocyte, an about 10-fold amount or more of an ACK-hemolysis medium was added to the pellet, and the mixture was maintained on ice for 15 to θ 20 minutes, and centrifuged at 1500 rpm at 4°C for 5 minutes. The pellet was washed twice with about 40 ml of a phosphate buffered physiological saline. The supernatant was aspirated, and

the resulting pellet was suspended in a cell banker at 106 to 107 cells/vial, and frozen and stored at -80°C or in liquid nitrogen.

Please replace Table 1 in its entirety with the new Table 1 below:

No.	9	8	7	6	5	4	3	2	1
Dilution Ratio (%)	100	10	1	0.1	0.01	0.001	0.0001	0	H ₂ O
Monkey DNA(μl)	100	10	10	10	10	10	10	_	_
Sheep DNA(μl)	_	90	-						
Total amount (µl)	90	90	90	90	90	90	90	90	_

<u>No.</u>	9	<u>8</u>	7	<u>6</u>	<u>5</u>	<u>4</u>	<u>3</u>	2	1
<u>Dilution Ratio</u> (%)	<u>100</u>	<u>10</u>	1	<u>0.1</u>	<u>0.01</u>	0.001	0.0001	<u>0</u>	<u>H₂O</u>
Monkey DNA(μl)	100	<u>10</u>	<u>1</u>	0.1	0.01	0.001	0.0001	=	
Sheep DNA(µl)	=	<u>90</u>	<u>99</u>	<u>99.9</u>	<u>99.99</u>	<u>99.999</u>	<u>99.9999</u>	<u>100</u>	=
Total amount (µl)	<u>100</u>	<u>100</u>	<u>100</u>	<u>100</u>	<u>100</u>	<u>100</u>	<u>100</u>	100	

Please amend the paragraph beginning at line 7, through line 18 as indicated below:

Here, as a control, for an amount corresponding to 250 ng of each of the following DNA:

only a sheep DNA (in Fig.3, lane 4),

a mixture of 0.0001% monkey DNA and 99.9999% sheep DNA (in Fig.3, lane 5),

a mixture of 0.001% monkey DNA and 99.999% sheep DNA (in Fig.3, lane 6),

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a mixture of 0.01% monkey DNA and 99.99% sheep DNA (in Fig.3, lane 7), a mixture of 0.1% monkey DNA and 99.9% sheep DNA (in Fig.3, lane 8), a mixture of 1% monkey DNA and 99.9% 99% sheep DNA (in Fig.3, lane 9), a mixture of 10% monkey DNA and 90% sheep DNA (in Fig.3, lane 10), and a monkey DNA alone (in Fig.3, lane 11),

PCR was performed simultaneously under the same conditions as for the specimen sample, and the product was used in electrophoresis.

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